Amendments to the Claims:

Please amend claims 3, 4, 7, and 8 as shown below.

This listing of claims replaces all prior versions and listings of claims in the application:

- (original) DNA material comprising either a T7 promoter or the xylA promoter, and a ribosome binding site from a Gram-postivie bacterium, and a reporter gene, which is operably linked to the promoter.
- 2. (original) DNA material of claim 1, wherein the reporter gene is a luciferase.
- 3. (amended) DNA material of any of claims claim 1 or 2, wherein the plasmid additionally comprises a selection marker and/or an origin of replication.
- 4. (amended) DNA material of any of claims claim1 to 3, wherein the DNA material comprises a sequence selected from the group comprising
 - (i) the sequence of (SEQ ID NO:5);
 - (ii) a sequence at least 90% identical to (SEQ ID NO:5);
 - (iii) the sequence of (SEQ ID NO:6); and
 - (iv) a sequence at least 90% identical to (SEQ ID NO:6)
- 5. (original) A method to determine whether a test substance has anti-microbial activity against Gram-positive bacteria, comprising the steps of
 - (i) incubating the test substance with bacterial cell extract of a Gram-positive bacterium and the DNA material of any of claims 1 to 4; and
 - (ii) detecting a signal resulting from the expression of said reporter gene.
- 6. (original) The method of claim 5 wherein the Gram-positive bacterium is Staphylococcus, Pneumococcus or Enterococcus.
- 7. (amended) The method of claim 5 or 6, wherein said bacterial cell extract is a bacterial S30 cell extract.
- 8. (amended) The method of claim 6-or 7 5, wherein said incubation is on a multiwell plate, suitable for use in a plate reader.